

Page I book references

EN/5738

Class, International, Electrical Pollution

Under reference (scientific works) Class, International Pollution, 1952.

657 b. Berlin, 1952, 2nd ed.

Contributors mentioned.

Persons: This book is intended for mathematicians, physicists, chemists, and
civil and mechanical engineers.

CONTENTS: The book consists of 59 papers by Russian specialists on problems in
mathematics and technology, particularly mathematics, physics, chemistry,
physics, civil and mechanical engineering. Some are given in English and
others in German and given at the end of each article. Some of the articles
are accompanied by references. No personalities are mentioned. At the back
of the book there are 25 references, all Russian.

NAME OF WORKER:

PART II. CONSTRUCTION

ARMED C. S. CHRISTIANSEN, INC. - NAME OF THE WORKER

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NAME III. MATERIALS

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AVRAM, C.N.; FILIMON, I.

Simple beton elements under the influence of eccentric compression.
Studii tehn Timisoara 9 no.3/4:325-337 Jl-D '62.

PANIN, N.; AVRAM, E.

New traces of vertebrates in the Miocene of the Rumanian sub-Carpathians. Studii cerc geol 7 no.3/4:455-484 '62.

AVRAM, G., MATEI, V.

New palaeontologic data on the Cretaceous Flysch in the northeastern part of the Retezat Valley Basin. Studii cerc geol geof geogr no. 2:321-327 '64.

I. Geologic Committee attached to the Romanian Council of Ministers. Submitted April 28, 1964.

MICHAELU, G., n. ad.; STEFANESCU, H.; AVRAM, E.; MATEI, V.; ANDREESCU, M.;
BUNAREANU, C.

Nature of the appearances of crystalline schists in the Zamurariu-
Prislop region. Studii cerc. geol. geogr. 9 no.2:511-518 '64.

1. Geological Committee and the Institute of Geology and
Laboratories. Submitted April 26, 1964.

ALIAN, G.; NICOLAE, I.

"Noramt Planning in Romania", p. 344, (REVISTA MURIER, Vol. 69, No. 8,
August 1984, Bucharest, Romania)

Sc: Monthly List of East European Accessories (EMAL), 1C, Vol. 4, No. 1,
March 1985, Uncl.

L 2045-66 EWA(d)/EWP(t)/EWP(k)/EWP(z)/EWP(b)/EWA(c) JD/EW
ACCESSION NR: AP5027064 RU/0017/65/000/004/0199/020; 23B

AUTHOR: Guida, O. (Engineer); Avram, Gh. (Engineer)

TITLE: Experiments concerning the manufacture of non-corrosive ferritic steel tubes in a mill for 3" seamless tubes of the stonksbank type

SOURCE: Metalurgia, no. 4, 1965, 199-203

TOPIC TAGS: ferritic steel, metal tube, metal test

ABSTRACT: On the basis of the tests reported in this paper, the authors conclude that non-corrosive ferritic steel H 17 could be used to advantage at the Republica Pipe Works, without major changes in manufacturing technology and with significant technical-economic advantages. Orig. art. has 6 figures and 7 graphs.

ASSOCIATION: Guida, Institutul de cercetari metalurgice (Metallurgical Research Institute); Avram, Uzinele de tevi "Repubica" ("Repubica" Pipe Works)

SUBMITTED: 00

ENCL: 00

SUB CODE: NH

NO RRF SOV: 004

OTHER: 004

JPRS

Card 1/1 00

AVRAM, I.

The AD-10 and the AD-12 ceramic fluxes for automatic welding; flux submitted and tested in the Lao Tze-jun Plants of Bucharest.

p. 52 (Metalurgie Si Constructii De Masini. Vol. 4, no. 4, Apr. 1957. Bucuresti, Romania)

Monthly Index of East European Publications (EEAJ) IC. Vol. 7, no. 2, February 1958

ZOSIN, C., prof.; MANESCU, N., dr.; AVRAM, J., dr.; MINZA, N., dr.; GHELERT, I. Lazar, ing.

A model of the artificial kidney made in the Medical Clinic III of the Medical Institute of Timisoara. Med. intern., Bucur 13 no.2: 297-302 F '61.

(KIDNEY, ARTIFICIAL)

MANESCU, N., dr.; AVRAM, J., dr.; METZ, A., dr.; HENNING, R., dr.;
POP, S., dr.

Nephrogenic anemia. Clinical and etiopathogenic considerations. Med.
intern. 14 no.9:1109-1116 S '62.

1. Lucrare efectuata in Clinica a III-a medicala (prof. C. Zosin),
I.M., Timisoara.

(KIDNEY DISEASES) (ANEMIA)

"APPROVED FOR RELEASE: 06/06/2000

CIA-RDP86-00513R000102620004-6

STEFANESCU, D., Prof.; TUDOR, N.; AVRAM, L.

Superficial tension of electrolyte solutions and their specific conductivity. Romanian M. Rev. 2 no.2:93 Apr-June 58.

(ELECTROLYTES)

superficial tension of electrolyte solutions & their specific conductivity)

APPROVED FOR RELEASE: 06/06/2000

CIA-RDP86-00513R000102620004-6"

Country : Romania
Subject : Organic Chemistry, Synthetic organic Chemistry

Ref. No.: Ref. Zhur-Khim., No.12, 1959, No.42274

Author :
Institut. :
Title :

Chem. Rev. :

Abstract : Substance (III) is hydrolyzed (PTC₂, 40° in ethyl acetate). This results in the formation of 'CH(CH₃)₂ having a melting point of 83°. The addition of Br to III produces RCBr-(CH₂Br)₂ (IV) which upon being heated is converted into RC(CH₂Br)=CHBr (V). In the presence of CH₃OH or of alcohol, IV is transformed into R(CH₂Br)₂R' where R'=CH₃ (VI) or C₂H₅ (VII). The synthesis of I is described according to the following arrangement: KCN+NH₄Cl+

Date: 2/7

Country: Romania G
Category: Organic Chemistry. Synthetic Organic Chemistry

Pub. Date: Ref Zhur-Khimiya, No.12, 1959, No.42074

Author: :
Institute: :
Title: :

Chem. Russ.: :

Matter of: ether). Unpurified IV (from 0.02 moles of III) is heated in vacuum for 1 hour at 105-110°; the melt is dissolved in hot CH₃OH and a 65% yield of V is separated; the melting point is 171°. CH₃OH is added to unpurified IV (from 0.04 mole of III); the mixture is boiled until the precipitation of HBr stops; a 59% yield of VI is separated; its melting point is 120-121° (from CH₃OH). VII is obtained in a similar manner by the action of alcohol; the melting

Card: 477

Country : Romania
 Category : Organic Chemistry. Synthetic Organic Chemistry G
 Abs. Jour. : Ref Zhur-Khimiya, No.12, 1959, No.42374
 Author :
 Institut. :
 Title :
 Orig. Pub. :
 Abstract : is separated. 1 mole of VIII is added to a solution of 1 mole of phthalic anhydride (or 166 g of $\text{C}_6\text{H}_4(\text{COCH}_3)_2$) and 2 moles of KOH in 400 ml of water; the mixture is heated, while evaporating the water, to 180-190°. This temperature is maintained for 30 minutes; upon cooling the mass is pulverized into powder which is washed with 10^3 NaHCO_3 ; the filtrate is acidified with concentrated HCl, and a 5% yield of IX is separated off; the melting point is
 Date: 6/7

Country : Romania
 Category : Organic Chemistry. Synthetic Organic Chemistry G
 Abs. Jour. : Ref Zhur-Khimiya, No.12, 1959, No.42374
 Author :
 Institut. :
 Title :
 Orig. Pub. :
 Abstract : 154° (from benzoyl). 0.36 moles of IX and 0.66 moles of SOCl_2 are heated at 50° for 2 hours; the excess SOCl_2 is distilled off in vacuum, and the unpurified I is separated; its melting point is 82°. The CO and HCl are separated by heating the unpurified X to 180-190°; the residue is distilled at 170-180°/10 mm; I is then obtained, its yield is 47%, melting point 105° (from CCl_4).
 -- V. Skorodumov

Artem, Marguerite

Chem

The catalytic isomerization of saturated hydrocarbons.
The mechanism of activation of aluminum chloride by water.
Costin D., Neftulescu, Marguerite, Avram, and Rivire Sham
(Acad. Rep. Populare Romane, Bucarest). Bull. Soc.
Chim. France 1953, 1225-72. In the isomerization of satd.
hydrocarbons by means of AlCl_3 , the catalyst must be acti-
vated either by an alkyl halide, which may be formed *in situ*
from a hydrogen halide and trace of olefins present in the
hydrocarbon feedstock, or by water. A mechanism for the
isomerization is proposed and supported. The hydrocar-
bons (2 mols), purified by treatment with H_2SO_4 followed
by hydrogenation over Ni and distn. over Na, were brought
into contact with 0.4 mole of AlCl_3 sublimed *in vacuo* and
activated with 0.4 mole H_2O , in a stream of CO_2 . The gas
produced was analyzed for H by combustion after removal
of the CO_2 in 50% KOH and of org. gases with active C.
Blanks were run on the CO_2 and on the system without the
activator. A trial with cyclohexyl chloride as activator for
 AlCl_3 in the isomerization of cyclohexane produced no H.
Vera I. Vivian

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0

AVRAM, II.

NENITESCU, C.; AVRAM, M.; SLIAM, E. Reduction of 5-nitouracil by sodium dithionite.

Studii cercetari chimice
Vol. 4, No. 1/2, Jan/June 1956

Bucuresti, Romania

SO: Monthly List of East European Accessions, (EEAL), IC, Vol. 5, No. 10,
Oct. 1956

"APPROVED FOR RELEASE: 06/06/2000 CIA-RDP86-00513R000102620004-6

AVROUIN, Marcelle

APPROVED FOR RELEASE: 06/06/2000 CIA-RDP86-00513R000102620004-6"

Cyclobutane series. I. 1,3-Disubstituted cyclobutane derivatives. Margareta Avram, Costin D. Neagu, and Maria Maxim (Akad. R.V.R., Bucharest, Romania). *Chem. Ber.* **80**, 1424-32 (1957); cf. *C.A.* **54**, 4512a. — The synthesis of 1,3-diaminocyclobutane (I) as well as several intermediate products and derivs. was described. Epibromohydrin (240 g.) and 290 g. PhCH₂Br heated 8 hrs. at 185-190° (internal temp.) with 0.2 g. HgCl₂ and the product distd. gave 243 g. crude PhCH₂OCH(CH₂Br)₂ (II), b.p. 148-55°; the forerun (b.p. to 160°) heated again with HgCl₂ and the product distd. gave 64 g. II, b.p. 148-55°; on repeated distn. II b.p. 148°. Epichlorohydrin (440 g.) and 615 g. PhCH₂Br treated with 0.6 g. HgCl₂, heated slowly (during 6 hrs.) to 165°, kept 6 hrs. at 165-170°, and 1 distd. gave 608 g. PhCH₂OCH(CH₂Br)CH₂Cl (III), b.p. 148-65°; from the forerun an addnl. 102 g. III was obtained; redistd. III b.p. 140°. Na (23 g.) was dissolved in 400 cc. EtOH, half the soln. and 88 g. CH₃(CO₂Et)₂, 400 cc. dry C₆H₆, and 1.64 g. crude II (or 132 g. crude III) heated 1.5 hrs. at 170° (bath temp.) in a 1.5 l. autoclave (pressure 0 atm.), cooled (neutral soln.), 400 cc. dry C₆H₆ and the remaining NaCOEt soln. added, heated 6 hrs. (with III 9 hrs.) at 170°, the cooled neutral soln. filtered from KBr (or KCl), and the filtrate concd. on a H₂O bath; the KBr was dissolved in H₂O, the soln. extd. with Et₂O, the ext. concd., the residue combined with the main fraction, and repeatedly distd.

to give 65 g. crude CH₃CH(OCH₂Ph)CH₂C(CO₂Et)₂ (IV), b.p. 178-33°, on redistn. b.p. 178°, d₄ 1.0083, n_D²⁵ 1.4810. Crude IV (160 g.) and 170 g. KOH in 420 cc. EtOH and 100 cc. H₂O refluxed 2 hrs. on a H₂O bath, evapd. *in vacuo* almost to dryness, the residue dissolved in 400 cc. H₂O, the soln. extd. with Et₂O, acidified with concd. HCl (voln. V), and the ppt. filtered off gave CH₃CH(OCH₂Ph)CH₂C(CO₂H)₂, m.p. 158° (H₂O). Soln. V prep'd. as above extd.

with 5 100-cc. portions Et₂O, the ext. dried, evapd., the residue heated at 10 min. until CO₂ evolution ceased, and distd. gave 98 g. CH₃CH(OCH₂Ph)CH₂CHCO₂H (VI), b.p. 194°, d₄ 1.13097. VI (2 g.) and 1 g. urea heated 3 hrs. at 150-155° gave the amide, m.p. 120° (H₂O). VI (98 g.), 60 cc. abs. EtOH, 80 cc. C₆H₆, and 2 drops H₂SO₄ refluxed 8 hrs. (H₂O separator used) gave 100 g. VI Et ester (VII), b.p. 130°, b.p. 175°, d₄ 1.0032, n_D²⁵ 1.5019, MR 65.00. VII (117 g.) in 800 cc. EtOH mixed with 0.5 g. Pd black, shaken with H at atm. pressure, after absorption of the catal. amt. H (8-8 hrs.) the catalyst filtered off, the filtrate concd., and the residue distd. gave 65 g. CH₃CH(OH).

CH₃CHCO₂Et (VIII), b.p. 110°, d₄ 1.0727, n_D²⁵ 1.4504, MR 36.13. VIII (30.4 g.) and 35.2 g. PhSO₃Cl cooled in ice H₂O treated portionwise with 20 cc. dry C₆H₆N, the mixt. kept 20 hrs. in an ice box, treated with 30 cc. H₂O, extd. with Et₂O, the Et₂O ext. washed successively with H₂O, dil. HCl, aq. Na₂CO₃, and H₂O, dried, and evapd. *in vacuo* (60° bath temp.) gave 55 g. PhSO₃ ester (IX) of VIII, purity 99.7%. IX (55 g.) diid. with 55 cc. EtOH, treated with 24 g. NaN₃ in 25 cc. H₂O, the mixt. boiled 10 hrs., cooled, the sep'l. PhSO₃Na dissolved by adding H₂O, the soln. extd. with Et₂O, the ext. washed with H₂O, and dried gave 27 g.

CH₃CHN₃CH₂CHCO₂Et (X), b.p. 94°, d₄ 1.0902, n_D²⁵ 1.4007, MR 42.64. X (25.6 g.) and 15 cc. NaH, H₂O (XI) heated 30 min. on a boiling H₂O bath, the excess XI evapd. *in vacuo* on the H₂O bath, and the residue cooled gave 23 g. hydrazide (XII), m.p. 70° (C₆H₆, ligroine). XII (23 g.) dissolved in 270 cc. cold (0°) N HCl, mixed with 160 cc. Et₂O, the mixt. treated dropwise at 0-5° with aq. NaNO₂

(10 g. $N_2N'D_2$ in 50 cc. H_2O) until free HNO_3 formed (detectable with KI-starch paper) the aq. phase sept., extd. with 2 50-cc. portions Et_2O , the ext. dried by stirring at 0° with $CaCl_2$, filtered, the Et_2O filtrate mixed with 10 cc. $PhCH_2OH$, the Et_2O slowly (2 hrs.) distd. (N evolution began at 20°), after the removal of the Et_2O the mixt. heated 1 hr. at 80° (N evolution ceased), and the excess $PhCH_2OH$ distd.

at 120-30° (oil bath temp.)/5 min. gave 24 g. $CH_3CH(NH_2)CH_2CH_2Ph$ (XIII), m. 60° (ligroine).

$CH_3CH(NHCO_2CH_2Ph)$ (XIII), m. 60° (ligroine). XIII (12.3 g.) in 100 cc. $BtOH$ and 0.2 g. Pd black shaken at 30-8° in an H atm. [the H often removed until it contained no more CO_2 (6 hrs.)] and the mixt. worked up as usual gave 2.3 g. I, b.p. 90°; dipelate m. above 250° ($BtOH$); hexachloroplatinate, crystals from H_2O . I (1.72 g.) heated 10 hrs. at 100° with 13.8 g. 95% HCO_2H and 10 g. 35% aq. CH_3O_2 the mixt. made alk. with 40% aq. $NaOH$, extd. with Et_2O in a Kutscher-Steudel app., and the product distd. gave 0.4 g. boron, b.p. to 70°, and 1.8 g. N,N,N',N' -tetra-Me deriv. (XIV) of I, b.p. 85° (dipelate m. above 240°). XV (3.8 g.) in 10 cc. abs. Et_2O treated with 7.6 cc. MeI and after 6 hrs. the product completely pp'd. with Et_2O gave XIV di-MeI salt, m. 249-50° ($MeOH-H_2O$); dipelate m. above 250°. VII (10 g.) and 20 cc. 80% XI heated 10 min. on the H_2O bath until homogenized, allowed to stand 2 hrs. at room temp., most of the liquid removed in *vacuo*, the residue treated with 10 cc. XI, and allowed to crystallize (12 hrs.) gave 13 g. hydrazide (XV) of VI, m. 100° ($BtOH$). XV (11 g.) dissolved in 100 cc. N HCl, treated dropwise with stirring with a concd. soln. of 3.6 g. $NaNO_2$ at 0-5°, the oily azide extd. with 150 cc. cold (0°) Et_2O , the ext. dried at 0° (0.25 hr.), treated with 150 cc. abs. $BtOH$, the Et_2O evapd. with a downward condenser (N evolution started at 60-60°), finally 50 cc. $BtOH$ added, the soln. refluxed 1 hr., and the $BtOH$ removed in *vacuo*.

gave 10 g. $CH_3CH(OCH_2Ph)_2CH_2CHNHCO_2Et$ (XVI), m. 65° ($BtOH$ or ligroine). XVI (0 g.) and 25 g. KOH in 70 cc. $MeOH$ refluxed 1 hr., the $MeOH$ distd., the residue mixed with H_2O , the mixt. extd. with Et_2O , the ext. washed with a

little H_2O , dried, and evapd. gave 6 g. $CH_3CH(OCH_2Ph)_2$.

$CH_3CH(NH_2)_2$ (XVII), b.p. 118°, n_D^{25} 1.52935, *M.R.* 53.0. XVII (8 g.) and 16 cc. Ac_2O boiled a short time and cooled gave the Ac deriv., m. 95° (ligroine); the Ac_2O filtrate concd. and the residue distd. gave an unidentified compd., b.p. 210°, m. 78° (ligroine). Hydrogenation of XVII with Pd black failed because of balling of the catalyst. To a complex of 59 g. CrO_3 in 300 cc. C_6H_6 prep'd. at 0° was added portionwise 28 g. VIII in 280 cc. C_6H_6 keeping the temp. below 20°, the mixt. kept 12 hrs. at room temp. and 1 hr. at 45°, dild. with ice H_2O , and extd. with 1:1 Et_2O -

C_6H_6 gave 18.1 g. $CH_3COCH_2CH_2CO_2Et$, b.p. 00°; semi-carbazone m. 205° (H_2O); 84% oxime (XVIII), b.p. 145 0°. XVIII (6 g.) in 500 cc. $BtOH$ hydrogenated with 0.5 g.

PtO_2 at 30-5° under normal pressure gave 1.7 g. $CH_3CH(NH_2)_2CH_2CH_2CO_2Et$ (XIX), b.p. 88°, and 2.4 g. fraction (apparently secondary amine), b.p. 180-5°, not further investigated. XIX gave an *N*-Bz deriv. (XX), m. 104.5-5.0° (ligroine), an *N*-phthalimid-1-cyclobutanecarboxylic acid, m. 178-80° (H_2O). XX treated with XI and the resulting

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hydrazide subjected to a Curtius degradation gave $\text{CH}_3\text{CH}(\text{NHCO}_2\text{Et})_2$

($\text{NHBa}_2\text{CH}_3\text{CHNHCO}_2\text{Et}$, m. 185° (aq. EtOH). IV re-

duced like VII gave 85% $\text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{C}(\text{CO}_2\text{Et})_2$
(XXI), b_d 120–2°, d₄ 1.1139, n_D 1.4488, M/R 52.17. XXI
oxidized like VIII gave 73% oxp diester, b_d 112–13°; semi-
carbazine m. 170°; 78% oxime (XXII), b_d 127°, d₄ 1.1583,
n_D 1.40885, M/R 85.03. XXII (12 g.) catalytically reduced
like XVIII gave 4 g. amino diester, b_d 98°, and 0 g. compd.,
b_d 195°, apparently the secondary amine. IX (10 g.) in

10 cc. EtOH treated with 10 g. KI in 5 cc. H₂O, boiled 10

hrs. on a H₂O bath, and worked up gave 5 g. $\text{CH}_3\text{CH}(\text{NHCO}_2\text{Et})_2$

(XXIII), b_d 113° (decompn.). XXIII (6
g.) in 100 cc. 80% EtOH treated portionwise during 12 hrs.
with 160 g. 3% Na-Hg with stirring, the supernatant liquid
sepd. from Hg, boiled 6 hrs., dried, and extd. with Et₂O

gave 1.0 g. $\text{CH}_3\text{CH}_2\text{CH}_2\text{CHCO}_2\text{H}$, b_m 110°, b. 194°;
S-benzylthiuronium salt m. 170° (EtOH). A more con-
venient method of prep. I from XII was as follows. XII
(31 g.) was converted, like in the prepn. of XIII but replac-
ing the PhCH₂OH by 100 cc. abs. EtOH, into 32 g. crude

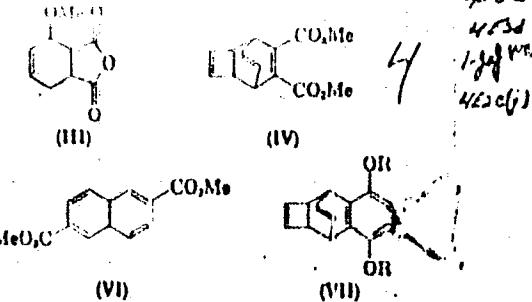
$\text{CH}_3\text{CHN}_2\text{CH}_2\text{CHNHCO}_2\text{Et}$ (XXIV). Crude XXIV (32

g.) subjected to a Curtius degradation gave 13 g. $\text{CH}_3\text{CH}(\text{NHCO}_2\text{Et})_2$

($\text{NHBa}_2\text{CH}_3\text{CHNHCO}_2\text{Et}$, b_m 18°, hydrogenated with Pd black
to 60% I, b. 166° [1,3-di-*t*-*c* deriv. m. 218° (EtOH)].

AVRAM, M

Investigations in the cyclobutane series. II. Cyclobutadiene.¹ Margareta Avram, Costin, D. Nitaescu, and Elie Iaica (Acad. R. V. R. Bucarest, Romania). Chem. Ber. 90, 1851-08 (1957); cf. C.A. 53, 10043f. Hofmann degradation of 1,3-bis(dimethylamino)cyclobutane as well as thermal fission of Reppe adducts from cyclooctatetraene and several dieneophiles result only in formation of butadiene (I) instead of the expected cyclobutadiene. A soln. of 1.8 g. 1,3-bis(dimethylamino)cyclobutane-2MeI (II) in 20 ml. MeOH is shaken 1 hr. with AgNO_3 (from 6 g. AgNO_3 and 10% aq. NaOH). After filtration and washing of the ppt. with MeOH, the filtrate is evapd. at 20-30° *in vacuo*. From a bath temp. of 120° up to 200° decompn. with formation of 20 ml. gas, which is caught over 60% KOH, takes place; the gas has a C:H ratio of 1:1.5. II (4.26 g.) is heated with 5 g. KOH in 5 ml. water for 1.5 hrs.; the resulting gas is led into a soln. of Et in CH_2Cl_2 . After evapn. a small quantity 1,2,3,4-tetrabromobutane, m. 117°, is left. II (4.26 g.) after treatment according to the first procedure, on distn. of the filtrate, gives 2 ml. MeOH, which on diln. with water yields an oil. After extn. with ether, drying, and evapg., the yellow oil (50 mg.) is mixed with 50 mg. maleic anhydride to give III, m.p. 97° (dioxane-ether) (Flalg, C.A. 44, 6808J) (MeO_2 ,



CCl_4) (28 g.) and 23 g. cyclooctatetraene is heated 8 hrs. at a bath temp. of 150-160°. At slightly higher temp. decompn. takes place to give 25 g. IV, b.p. 140-160°. On heating 0.1 mole (24 g.) IV 15 min. to about 300°, 360-420 ml. gas is vigorously evolved and proves to be I (16-17% yield). From the residue there is obtained 4-6 g. di-1,6-2,6-naphthalenedicarboxylate (V), m. 187°, d-6.6 g. (30-50) di-Me phthalate, b.p. 140-60°, and 6.6 g. VI, m. 120°. At 300° with 30% Pd-C VI gives V. When VII ($R = \text{Me}$), m. 105° (Reppe, et al., C.A. 43, 6194g), is heated to the b.p., I is obtained in 26% yield, and from the residue 60-64% 1,4-dimethoxy naphthalene, m. 88°, and 1,4-dimethoxyphenanthrene, m. 124.5-125°, are isolated and identified. VII ($R = \text{Ar}$), m. 136-7° (loc. cit.), gives on thermal degradation 1,4-diacetoxynaphthalene, m. 123-6°, and 1,4-diacetoxynaphthalene, m. 137°. Fission of the adduct of cyclooctatetraene and α -naphthoquinone, m. 160-2°, gives I, anthraquinone, and a mixt. of isomeric benzanthraquinones, m. 160° (MeOH) (the mixed m.p. with authentic 1,2-benzanthraquinone, m. 168°, is 155°).

C. B. Frating

Distr: 4E2c(j)/4E3d

1/ Cyclobutanes. III. Benzocyclobutadiene. Costin D.
Nemțescu, Măxineni Axram, and Radu Dinn (Tech.
Hochschule, Bucharest, Romania). *Chem. Ber.* 90, 2541-4
(1957); cf. *ibid.* 1957.—Formation of dihydrobenzodiphen-
ylene (I) from 1,2-dibromobenzocyclobutene (II) and Zn
(cf. Cava and Napier, *C.A.* 51, 11810l) suggests that
benzocyclobutadiene (III) is a distinct but short-lived inter-
mediate. The cyclopentadiene (IV) adduct of III is now
obtained by reaction of II with Li-Hg, then IV. Similar
reaction with furan under these conditions gives an amor-
phous polymer. A soln. of 80 g. *p*-C₆H₄(CHBr)₂ in 600
cc. Me₂CO (very dry) is refluxed 52 hrs. with 105 g. NaI in
100 cc. Me₂CO, filtered; the soln. evapd. to 750 cc., decolor-
ited with Na₂S₂O₃, and dild. to 11. with H₂O, the ppt. sepd.,
and the soln. extd. with Et₂O gives 50% II, m. 62-3°. A
soln. of 3 g. II in 60 cc. abs. Et₂O is shaken 10-12 hrs. with
100 g. 0.5% Li-Hg (cooling at first), sepl., washed with
Et₂O, and evapd. to give 80% I, m. 74°. A soln. of 18 g. II
in 25 cc. abs. Et₂O shaken with 600 g. 0.5% Li-Hg and 19 g.
IV gives 62% 1,4-endomethylene-1,4,11,12-tetrahydrodiphenyl-
ene (V), b₅ 82-4°, n_D²⁰ 1.57317, d₄²⁰ 1.0639. V absorbs 1.2
molecules H over Pd. V forms adducts with PhN₃ (m. 132°)
and CH₂N₂ (m. 10°). The latter heated to the b.p. in a
CO₂ stream forms pyrazine. M. A. Simkins

AVRAM, M.

Distr: 4E2e(j)/4E3d

Brominated derivatives of 2-phthalimidopropane. Costin
D. Nechita, M. Dragulescu, V. Ilie, and Maria Maxim.
Acta rep. populare Române, Stiinte Exacte si Chim. 6, 239-45
(1958). — α-Phthalimidobutyric acid (I) was prepd. from
K H phthalate and HO₂CCMe₂NH₂HCl. From the acid
chloride of 1,2-phthalimidopropane (II), m. 108° was ob-
tained; on bromination with N-bromo-N-chloro-N,N-dimethyl
1-bromo-2-phthalimidopropane (III), m. 98°. Brominating
III gives 1,1,3-tribromo-2-phthalimidopropane (IV), m.
101°. The 2 Br of IV is easily substituted by methoxy or
ethoxy group to yield, resp., 1,3-dibromo-3-methoxy-2-
phthalimidopropane, m. 117-18°, and 1,3-dibromo-2-
ethoxy-2-phthalimidopropane, m. 135°. J. Berall

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3 May

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AVRAM, MARGARETA

Distr: 4E3d/4E3b/4E2c(j)

Cyclobutane series. V. Some bromo derivatives of cyclobutanes. Margareta Avram, Elise Marin, and C. D. Nenitescu (Acad. R.P.R., Bucharest, Romania). *Acta Pol. pop. Romae, Studii cercetari chim.* 7, 155-17 (1959) (German summary); cf. *C.A.* 53, 16993f; preceding abstr. — Both *cis*- (I) and *trans*-1,2-cyclobutanedicarboxylic acid (II) were converted into a mixt. of *cis*- (III) and *trans*-1,2-dibromocyclobutane (IV) by the Hunsdiecker reaction (which was therefore not stereospecific). (38 g.), m. 139.5-140.5°, or its anhydride (V) in 180-200 cc. H_2O was added to 85 g. $AgNO_3$ in 1700 cc. H_2O . The Ag salt of I (80-85 g.) pptd. The salt was dissolved in 1100 cc. dry CCl_4 , stirred, and cooled to 0°. Br (87.0 g.) was added (the temp. rose to 65°), the soln. filtered, 39, eq. $NaHSO_3$ added. Till decoloration occurred, the soln. washed (H_2O), dried ($MgSO_4$), the solvent evapd. in a column, and the residue fractionated twice in *vacuo* to give a mixt. (10.2-21.0 g.) of 14% III and 78% IV. II, m. 131-1.5°, was prep'd. from the di-Et ester of I, b.p. 138°, by converting it with NaOBt to the di-Bt ester of II, b.p. 120°, and then boiling with 6N HCl 12 hrs. The Ag salt of II was converted into a mixt. of 26% III and 65% IV as above. 1-Bromocyclobutene was dissolved in 130 cc. Isopentane and dry HBr was introduced at 0° under ultraviolet illumination 3-4 hrs. Fractionation gave 10 g. III, 97% purity, b.p. 102°, n_D 1.45 to 1.42, d₄ 2.0103, n_D 1.54981. IV, 97% purity, b.p. 93-4°, m. -3.5 to -4.0°, d₄ 1.03051, n_D 1.43441, was prep'd. in $CHCl_3$ by bromination of cyclobutene. A mixt. (21.4 g.) of III and IV, 21.4 g. $AlBr_3$, and 0.036 cc. H_2O was stirred in 180 cc. CS_2 at 0-15° 10 hrs. The CS_2 soln. was decanted from the red resl which sepd., washed, evapd., and the residue fractionated in *vacuo*. Thus, 15-16 g. mixt. was obtained contg. 11% III, 47.6% IV, and 31.5% 1,3-dibromocyclobutane (VI) (detd. by gas chromatography). This mixt. (10.7 g.) and 20 g. powd.

KOH was heated at 100-105° 1 hr. From the gas evolved, 1.4 cc. $HC\equiv CCH:CH_3$ (VII) condensed in a trap, and $HC\equiv CCH:CH_3$ was adsorbed in a Cu(I) soln. VII was identified by converting it into tetrabromovinylacetylene. A mixt. (10.7 g.) of III and IV dissolved in 10 cc. CS_2 was mixed with 3 g. $AlBr_3$ dissolved in 80 cc. CS_2 at 0°, instd. with dry HBr, agitated 2 hrs. at 0° and 6 hrs. at 15°. The solvent was decanted, evapd., and the residue dried. *In vacuo*. Tribromobutane (VIII) (10 g.), b.p. 104-5°, d₄ 2.2028, n_D 1.68308, was obtained. VIII (2 g.) heated at 110° with 5 g. KOH gave VII. 1,2-Dibromocyclobutane-1,2-dicarboxylic acid (IX) (30.3 g.) was dissolved in 20 cc. H_2O , neutralized with NH_3 to pH 7, and dropped into a soln. of $AgNO_3$ (34.6 g. in 300 cc. H_2O). The Ag salt of IX (47.50 g.) pptd. The Hunsdiecker reaction of this (18.8 g.) gave 3 g. 1,1,2-tetrabromocyclobutane (X), m. 120° (MeOH), $AlBr_3$ (1.50 g.) dissolved in 100 cc. CS_2 was added to 18.6 g. X at 0° and agitated at 10-20° 10 hrs. A mixt. of isomers ($C_4H_7Br_3$) was obtained, b.p. 114-15°. I (7.2 g.) dissolved in 60 cc. C_6H_6 was mixed with 7.0 g. dry pyridine and 22.2 g. $Ph(OAc)_2$, and stirred at 50-60° 1 hr. Cyclobutene (XI) evolved was introduced into a soln. of Br in CH_2Cl_2 , from which 1.6 g. 1,2-dibromocyclobutane was obtained, b.p. 95-100°. A flask contg. 180 cc. dry CCl_4 , 17 g. bromosuccinimide, and 0.6 g. benzoyl peroxide was filled with carefully dried XI and agitated at 70-75° 0 hrs. The CCl_4 soln. was decanted from the polymer (oil) sepd., and the solvent evapd. By fractionating the residue, 1,2-dibromocyclobutane (3.8-3.5 g.), b.p. 80-85°, was obtained. VI. The decomposition of quaternary salts of 1,3-diaminocyclobutane. Margareta Avram and C. D. Nenitescu. *Ibid.* 109-74.—The attempted Hofmann degradation of 1,3-bis(dimethylamino)cyclobutane dimethobromide (I) in the presence of PhLi (II) was found to be anomalous, giving phenyleyclobutene (III). I (18 g.), m. above 205°,

Aurum, N,N'-dimethyl-N,N'-bis
and Pentacessic, C.C.

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was prep'd. from 9.6 g. 1,3-bis(dimethylamino)cyclobutane, (IV) dissolved in 10 cc. dr. MeOH at 0° by satg. it with dry MeBr and allowing the rxnt. to stand 12 hrs. To 100 cc. Et₂O soln. of II, obtained from 9.6 g. PhBr, 4.9 g. I was added. The soln. was agitated 80-100 hrs. Me₂N was isolated as its picrate (3 g.). Also obtained were a white cryst. product, m. 157° (EtOH), and 0.46 g. III, b. 80-8°. III was converted into 2,4-dinitrophenyl phenylcyclobutyl sulfide, m. 98°. IV (0.6 g.) was added to 15 cc. 30% H₂O₂ and stirred at 0-10° 10 hrs. to give 1,3-bis(dimethylamino)cyclobutane N,N'-dioxide, m. 85-90° (hygroscopic); picrate m. 181° (MeOH). T. Saito

Distr: 4E2c(j)/4E3b/4E3d

Preparation of cyclobutadiene. C. D. Nenitzescu, Margareta Avram, Riza Marcia, Maria Maxim, and Dumitru Dinu (Chem. research center, Acad. R.P.R., Bucharest, Romania). Acad. rep. populare Romane. Studii cercetari chim. 7, 481-504(1959). — A literature review (and results of exptl. work based on it) was presented in connection with the synthesis of cyclobutadiene or alkyl- and aryl-cyclobutadiene derivs. Hofmann degradation of the diquaternary base of 1,3-diaminocyclobutane, thermal decompn. of cyclooctatetraene adducts, and attempts to prepare cyclobutadiene from brominated derivs. of cyclobutane were performed. Cyclobutadiene was detected in AgNO₃ and Hg complexes, as well as (for a short period) in the gaseous state. Transformation of the Hg-cyclobutadiene complex into cyclooctatetraene was accomplished. Alkylated and arylated derivs. of cyclobutadiene were prepd. (32 references) (Summaries in Russian and English).

M. Lapidot

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1-BW(BW)
2-777(LNB)(way)
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Centralul de cercetari chimice al Academiei R.P.R.,
sectia de chimie organică, București.

AVRAM, Margareta; MARICA, Eliza; NENITZESCU, Costin D. [Nenitescu, C.D.],
acad.

Research in the series of cyclobutane. V. On the bromoderivatives
of cyclobutane. VI. Research in the obtainment of quaternary salts
derived from cyclobutanediamine. Rev chimie 4 no.2:253-270 '59.

(EEAI 9:?)

1. Aus dem Chemischen Forschungslaboratorium der Akademie der RVR,
Bukarest. 2. Relecteur en chef, Comite de redaction, Revue de
Chimie; Mitglied der Akademie der Rumänischen Volksrepublik (for
Nenitescu)

(Ring compounds) (Cyclobutane) (Bromine)
(Quaternary compounds) (Salts) (Cyclobutanediamine)

AVRAM, M.

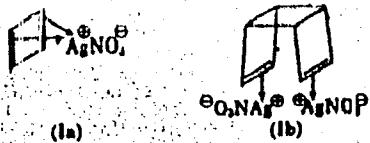
A review of Vols. 1-2 of C. D. Nonatescu's Tratat elementar de Chimie organică (Elementary Treatise on Organic Chemistry). p. 125.

REVISTA DE CHIMIE. Bucuresti, Rumania. Vol. 10, no. 2, Feb. 1959.

Monthly List of East European Accessions. (EEAI), LC. Vol. 8, no. 9, Sept. 1959.
Uncl.

AVRAM, I.

Studies in the cyclobutane series. IV. A silver saltate complex $C_4H_6AgNO_3$. Mihai Avram, Elia Mirica, and Cosmin Neagu (Acad. R.V.R., Bucharest, Romania). *Acta. Ber.* 92, 1088-91 (1960); *c.f. C.A.* 52, 18335g.—The elimination of Br from 1,2,3,4-tetrabromo-cyclobutene (I) with Li-Hg yielded an unstable Hg deriv. which with $AgNO_3$ gave a cryst. complex having the structure Ia or Ib. Cyclooctatetraene dibromide oxidized in Me_2CO with $KMnO_4$, at -5° by the method of Cope and Burg (*C.A.* 47, 3925) gave 43% 3,4-dibromocyclooctadien-1,2-dicarboxylic acid (II), m. $191-2^\circ$ (Et_2O -petr. ether). II



(41.2 g.) in 100 cc. H_2O adjusted with 3% aq. KOH to pH 7, treated dropwise with stirring with 40.5 $AgNO_3$ in 900 cc. H_2O , and the ppt. filtered off rapidly, washed with H_2O , Et_2NH , and Et_2O , and dried 24 hrs. in the dark over P_2O_5 gave 64 g. Ag salt (III) of II. III (40 g.) and 80 cc. CCl_4 refluxed with the slow removal of 30-40 cc. CCl_4 through the condenser, the residue cooled, treated with 8.1 cc. Br, heated briefly on the water bath, refluxed 15 min., cooled, filtered, washed, dried, and evapd., and the residue (9.16 g.) dissolved in dry Et_2O , kept 24 hrs. over K_2CO_3 , and evapd. gave I, b, $115-20^\circ$, m. 104° ($MeOH$ or petr. ether). I (1.37 g.) and 42 g. 0.5% Li-Hg shaken 15-20 hrs. at room temp. with 35 cc. abs. Et_2O , the Et_2O layer decanted, the Hg sludge washed with Et_2O , the combined exts. washed, dried, and evapd. under N gave a pale yellow oil with an acrid, characteristic odor, decomp. about 60° . The Et_2O soln.

from a similar run shaken 1-3 min. with satd. aq. $AgNO_3$ and filtered, and the residue (0.82 g.) washed with Et_2O and recrystd. from $EtOH$ gave Ia-Ib, m. 140° . Ia-Ib (0.22 g.) treated under CO_2 with dil. NH_4OH gave 0.3 cc. gas. Ia-Ib (0.53 g.) powdered and added gradually to Br in CH_2Cl_2 , and the mixt. filtered, washed, dried, and evapd. gave 0.1 g. oil, contg. 42.8% Br and 7.18% N, which decompd. on attempted distn. *in vacuo*. The formation of cyclooctadiene which existed for short periods of time in the gaseous state was demonstrated by treating Ia-Ib with a small amt. of steam and sweeping with air through Ascarite and $Mg(ClO_4)_2$ to give a gas which formed Ia-Ib with $AgNO_3$ soln.

F. W. Hoffmann

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Distr: 4E2c(j)/4E3d

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JA5(r10)Mn

Deaminations of 3-benzyloxy-1-cyclobutylamine by nitroso acid. Maria Maxim, Margareta Avram, and C. D. Nease, J. Am. Chem. Soc., 72, 1920, 1920.—3-Benzyloxycyclobutylamine (31.5 g.) was treated with 230 cc. 10% AcOH and a soln. of 21 g. NaNO₂ in 4.5 cc. H₂O (at -3° and with stirring). The mixt. was stirred 2 hrs. at 0°, kept overnight at room temp., and extd. with ether, the ext. neutralized with Na₂CO₃, the undesired resin decanted, the ext. dried (MgSO₄), and the Pt₄O evapd. to give 82% benzyl alc. (I), b.p. 93°, and 42.8% 3-benzyloxy-1-cyclobutanol (II), b.p. 145°, d₄ 1.073; n_D²⁰ 1.5249. It was identified as the phenylurethan, m.p. 78°, and II by its phenylurethan, m.p. 115° (EtOH). The overall yield of the diazotization was 74.3%. In this reaction, no formation of cyclopropane compds. was observed and the Denjanov rearrangement did not occur. To confirm the cyclobutane structure of II, the following reactions were carried out. II (8 g.) in 60 cc. anhyd. C₂H₅N was added (room temp., with stirring) to a complex of 10.5 g. C₂O₄ and 105 cc. anhyd. C₂H₅N, the mixt. stirred 0.5 hr., kept overnight, heated 2 hrs. at 0°, poured over ice H₂O, and filtered. The filtrate was extd. thrice with 1:1 ether-H₂O, the exts. washed with H₂O, dil. HCl, and H₂O again, dried (MgSO₄), filtered, and the solvent evapd. The residue gave 82% 3-henzyloxy-1-cyclobutanone (III), b.p. 153°, d₄ 1.0506, n_D²⁰ 1.6223; semicarbazone m.p. 101° (EtOH). To 1 g. III in 1.5 cc. C₂H₅N was added portionwise 0.6 g. KMnO₄ in 8 cc. C₂H₅N and 3.5 cc. H₂O, the mixt. left overnight, filtered, and concd. *in vacuo*. Addn. of a little H₂O and acidification with HCl gave BzOH, m.p. 121°. To an ice-cooled mixt. of 5 g. II and 4.8 g. PbSO₄Cl was added dropwise 2.6 cc. anhyd. C₂H₅N, the mixt. kept overnight, H₂O added, and the oil layer sep'd. The aq. layer was extd. with ether, the oil and ether layers washed with H₂O, dil. HCl and Na₂CO₃ soln., dried (MgSO₄), filtered, and evapd.

*gave 82% 3-benzyloxy-1-cyclobutanol benzenesulfonate (V). Finely powdered LiAlH₄ (1 g.) was added with stirring to 7.8 g. crude V in 80 cc. abs. ether, the mixt. stirred 0.8 hr., refluxed 1.4 hrs., and 2 cc. H₂O, 1.5 cc. 20% NaOH, and 7 cc. H₂O added with cooling and stirring. Filtering, washing with ether, drying the ether soln. (MgSO₄), filtering, evapg. the ether, and distg. gave (as a residue) 63% benzyl oxycyclobutane (VI), b.p. 98°, b.p. 80°, d₄ 0.9769, n_D²⁰ 1.5114, and 16% 3-benzyloxy-1-cyclobutanol, identified by its phenylurethan, m.p. 115°. Through a soln. of 0.6 g. VI in 7 cc. MeNO₂, dry HBr was passed for 5 min. After the soln. was cooled and kept 1 hr. at room temp., a part of the HBr was removed *in vacuo*, the soln. neutralized with Na₂CO₃ soln., and the org. layer dried (MgSO₄), filtered, and distd. The fraction, b.p. 100° [contg. MeNO₂ and cyclobutanol (VII)], was treated with PhIsocyanate to give cyclobutanol phenylurethan, m.p. 127° (peir. ether). Dry HBr gas was passed through 7.5 g. II in 30 cc. MeNO₂ for 5-10 min. and excess gas eliminated *in vacuo*. The soln. was neutralized with anhyd. K₂CO₃ and filtered, and the filtrate [contg. 1,3-dihydroxycyclobutane (VIII)] treated with PhIsocyanate to give (after 3-4 hrs.) 35.5% urethan, m.p. 207°. Another proof of the cyclobutane ring preservation, after deamination, was the prepn. of di-Et 3-hydroxycyclobutane-1,1-dicarboxylate (IX) by the diazotization of the di-Et 3-aminocyclobutane-1,1-dicarboxylate. The yield was 78.5% and its phenylurethan m.p. 115° (EtOH). Oxidn. of IX gave 85% di-Et 3-cyclobutane-1,1-dicarboxylate (X), b.p. 129°; semicarbazone m.p. 177° (EtOH). The reason for the stability of the cyclobutane ring in II and IX and of the absence of any cyclopropane ring may have been the presence of substituents with -I effect, and the fact that in these cases the cyclopropane ring could not be stabilized by resonance.* M. J. S.

IVANOV, MARGARETA

Distr: 483d

V. Investigations in the cyclooctane series. VII. Di-benzo-tricyclooctadiene and dibenzo-cyclooctatetraene. Margareta Avram, Delia Dinn, Georg Matzen, and Costin T. Nemecanu (Acad. R.V.R., Bucharest, Romania). *J. Am. Chem. Soc.* 93, 1789-1813 (1971); *C.A.* 54, 8604f. — Benzo-cyclobutadiene (I) is slow to react in the presence of Ni(CO), (II) to 3,4;7,8-dibenzo-tricyclo[4.3.0.0^{1,4}]octa-3,7-diene (III). III on warming was converted, with cleavage of the middle cyclobutane ring, to 1,2;5,6-dibenzo-cyclooctatetraene (IV), which, with Br, underwent ring contraction to 2,5-dibromo-3,4;7,8-dibenzo-cyclo[4.2.0]octa-3,7-diene (V). 1,2-Di-Br deriv. (20 g.) of I and 8.5 g. II in 310 cc. dry Et₂O shaken 0.5 hr manually and 3 hrs. mechanically with 600 g. 0.6% Li-Hg (or 1800 g. 0.5% Na-Hg), the Et₂O layer decanted, the Hg sludge washed with Et₂O, the combined Et₂O solns. washed with H₂O and filtered from 7.6% yellow solid contg. 16% N, and the filtrate worked up yielded 7.8 g. III, m. 133° (hydrogen or Et₂O). III (1 g.) in 20 cc. α -CaH₅Cl, refluxed 4–5 hrs. under N and evap'd. *In toto* gave 0.73 g. IV, m. 109° (sublimed *in vacuo*). IV in EtOH oxidized and treated with H₂O₂ gave phthalic acid and phthalic anhydride. IV (0.1 g.) in 2 cc. MeOH treated with satd. aq. AgNO₃ gave IV·AgNO₃, m. 222° (EtOH). IV (0.1 g.) in a little C₆H₆ treated with PhCN-Pd(3% C₆H₆) gave IV·PdCl₄, m. 224°. IV (1 g.) in 3 cc. C₆H₆

treated slowly with stirring at –5 to 0° with 1.5 g. Br in CH₂Cl₂, kept 0.5 hr. In the cold, and evap'd. *in toto* yielded 1.72 g. V, m. 157° (hydrogen, b. 80–80°). V (350 mg.) in 180 cc. dry Et₂O shaken 0.5 hr. with 10 g. 0.6% Li-Hg, filtered under N from the Hg sludge, washed with H₂O, and worked up yielded 90–120 mg. 4,5-benzo-tricyclo[4.4.0.0^{1,4}]dodeca-1,4,7,9,11-pentaene (VI), m. 78–80°; upon recrystalln. the m.p. rose and the O-content of the material increased. VI kept 12 hrs. in air or treated several hrs. with an air jet in Et₂O soln. gave the peroxide (VII), m. 135° (decompn.). VII hydrogenated over Pd-C gave a hydrocarbon, m. 103°, contg. 91.07% C and 8.06% H. VI (90 mg.) and 80 mg. maleic anhydride heated 5 min. at 80–90° and extd. with Et₂O gave 30 mg. hygroscopic adduct, m. 250–5°. VI and N-phenylmaleimide gave (similarly) an adduct, m. above 250° (decompn.). V (2 g.) in 8 cc. glacial AcOH and 1.09 g. KOAc refluxed 12 hrs., filtered, and dried with H₂O, and the product isolated with Et₂O yielded 1.3 g. 2,5-di-AcO analog (VIII) of V, m. 192° (MeOH). VIII (0.13 g.), 0.112 g. KOH, and 5 cc. EtOH refluxed 2 hrs. gave 0.0778 g. 3,4;7,8-dibenzo-cyclo[4.2.0]octa-3,7-diene, m. 133°; the mother liquor (conc'd.) gave orange-yellow crystals, C₁₄H₁₀, m. 133° (MeOH and EtOH). The infrared absorption spectra of these compds. were discussed.

J. W. Hoffmann

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1-JAC(8W)
1-JAC(17B)
1-JAC(18A)

MATEESCU, Gheorghe; AVRAM, Margareta; DINU, Doina; NENITESCU, Costin D., acad.

Infrared spectrum of dibenzotetra-cycloadiene. Studii cerc chim 9
no. 3: 427-434 '61.

1. Institutul de chimie al Academiei R.P.R., Sectia de chimie organică,
Bucuresti, 2. Redactor responsabil "Studii si cercetari de chimie"
(for Nenitescu).

AVRAM, Margareta; MATIESCU, Gheorghe D.; DINU, Doina; DINULESCU, Ilie G.;
NENITESCU, Costin D., acad.

Studies in cyclobutene series (VIII). Adducts of benzocyclobutadiene
with some cyclic dienes. Studii cerc chim 9 no.3:435-445 '61.

1. Institutul de chimie al Academie R.P.R., Sectia de chimie organica,
Bucuresti.

MATEESCU, Gh.D.; AVRAM, Margareta; NENITESCU, C.D., acad

Infrared spectra of some additional products of cyclooctatetraene with different phyledienes. Studii cer chim 10 no.1:65-72 '62.

1. Laboratorul de chimie organica al Institutului Politehnic si Centrul de cercetari speciale al Ministerului Sanatatii, Bucuresti.
2. Membru al Comitetului de redactie si redactor responsabil, "Studii si cercetari de chimie" (for Nenitescu).

AVRAM, Margareta; STERNBERG, Renée; DINULESCU, I.G.; NENITESCU, C.D., acad.

Condensation of 1,3-diiod-2-phenylpropane with ethyl malonate. Studii
cer chim 10 no.1:73-80 '62.

1. Centrul de cercetari chimice al Academiei R.P.R., Sectia de chimie
organica, Bucuresti. 2. Membru al Comitetului de redactie si redactor
responsabil, "Studii si cercetari de chimie" (for Nenitescu).

MATEESCU, G.D.; AVRAM, Margareta; DINU, Doina; NENITESCU, C.D.
[Nenitescu, C.D.]

Infrared spectrum of dibenzotricyclooctadiene. Rev chimie 8
no.1:13-20 '63.

1. Institute of Chemistry of the Academy of the R.P.R. Bucharest.
2. Member of the Academy of the R.P.R. (for Nenitescu).

AVRAM, Margareta; MATEESCU, G.D.; DINU, Doina; DINULESCU, I.G.;
NENITZESCU, C.D. [Nenitescu, C.D.] Member of the Academy of the R.P.R.

Investigations in the cyclobutane series (VIII). Rev chimie 8
no.1:77-86 '63.

1. Institute of Chemistry of the Academy of the R.P.R., Bucharest.

NENITESCU, C.D., academician; AVRAM, Margareta; POGANY, I.I.; MATEESCU, Gh.D.
FARCASIU, Malvina.

Synthesis and thermal decomposition of tricyclic ..
[4.2.2.0^{2.5}]-deca-3,7,9-triene. Studii cerc chim II no.1:
7-18 '63.

1. Sectia de chimie organica a Centrului de cercetari chimice
al Academiei R.P.R., Bucuresti.

DINULESCU, Ilie G.; AVRAM, Margaret; MARINESCU, Margareta; MIHAILOIU,
Costin D.

Research in the cyclobutane series. Pt.14. Studii cercet. chim. 12
no.5:357-364 '64

1. Center of Organic Chemistry Research, Romanian Academy, Polity
St., no.1, Bucharest.

TURGU, T., prof.; PAINA, N., dr.; IOHESCU, Gabriela, dr.; DUTU, Doina, dr.;
AVRAM, Maria, dr.; FLORESCU, O., dr.

The pronounced increase in rats infected with the causal agents
of anthropozoonoses demands intensification of rat eradication.
J. hyg. epidem. (Praha) 9 no.1:75-76 Ja-F'64

Hartman, Nicolina

Synthesis and properties. The preparation of phenyl-1,3-dimethyl-4-nitro-5-oxo-1,2-dihydro-1H-pyrazole.

Journal of the American Chemical Society, 1942, Vol. 64, No. 10, p. 4032.
An 0.032 g. benzylidenebenzaldehyde was dissolved in 0.5 g. Na in 10 ml. MeOH and 0.05 g. freshly dried zinc dust was added from reagent bottle. After 5 hr. in 25-30 ml. warm H₂O, cooling, and separating the product with acid to pH 4.0 gave a dusty white solid which, washed with H₂O and recrystallized from 10-20 ml. hot EtOH, gave 0.032 g. Phenyl-1,3-dimethyl-4-nitro-5-oxo-1,2-dihydro-1H-pyrazole, m.p. 113°.

Gary Geissel

Cambridge

"APPROVED FOR RELEASE: 06/06/2000

CIA-RDP86-00513R000102620004-6

AVRAM, N. Constantin; FILIMON, I.; FRIEDRICH, R.

Calculation of reinforced concrete frames in the plastic stage. Bul St si Tehn Tim 8 no.1:161-167 Ja-Je '63.

APPROVED FOR RELEASE: 06/06/2000

CIA-RDP86-00513R000102620004-6"

"APPROVED FOR RELEASE: 06/06/2000

CIA-RDP86-00513R000102620004-6

AVRAM, N. Constantin; VOLVA, N.I.; PAUNESCU, M.; MIHAESCU, A.; CRUIA, A.

New results of revibrating concrete. Bul St si Tehn Tim 8
no.1:169-185 Jan '63.

APPROVED FOR RELEASE: 06/06/2000

CIA-RDP86-00513R000102620004-6"

RUMANIA / Soil Science. Mineral Fertilizers.

J-4

Abs Jour: Ref Zhur-Biol., No 8, 1958, 34389.

Abstract: under fall plowing and as side-dressing in the presence of 6 leaves, N 20 and after the closing of rows N 20 and K 60. The doses indicated showed good results at many experimental stations in various territories of the land. -- V. V. Prok-hoshev.

Card 2/2

M

Country : RUMANIA

Category: Cultivated Plants. Grains.

Its Jour: RZhBiol., No 11, 1958, No 48880

Author : Davidescu, D.; Birsan, N.; Avram, P.; Enescu, D.; Popescu, I.; Vanea, I.; Davidescu, E.

Inst : Sci. Res. Inst. of Agriculture.

Title : Results of the Experiments on the Introduction of Fertilizers Under Corn, and Spring and Winter Wheat.

Orig Pub: An Inst. cercetari agron., 1957, 24, No 5, 47-69

Abstract: Experiments on the application of fertilizers were conducted during 1953-1955 at the following stations: at the Experimental Station of Merkuleshti in Konstantsa region, on applying basic fertilizers under corn; at the experimental stations of Kypic Turziy

Card : 1/2

M

RZhBiol., No 11, 1958, No 48880

in Kluga region and of Bucharest; at the Suchava region, on plowing potassiumite from Mereshti under spring wheat; at the experimental stations of Merkuleshti and Studin in Krivovskaya region and Moara Domnyaska in the Bucharest region, on applying side-dressing to winter wheat according to the vegetation phases. The article indicates the most suitable doses of the basic fertilizers under the corn (application of 30 kg/ha of manure and 20 kg/ha of manure plus N 40 P 30 K 20), under the spring wheat (P_{2O_5} 40 kg/ha) and winter wheat (N 30 kg/ha). -- A.F. Khlystova

Card : 2/2

Avram, P.

HIROX Oh
BUKAVU (in case); given name

Country: Romania

Academic Degree: --

Affiliation: --

Sources: Bucharest, Comunicarele Academiei Republicii Populare Romane,
No 4, 1961, pp 473-478.
Data: "The Efficiency as Fertilizer of the Manganese Residue of
Isobutyl Succinic Acid."

Co-authors:
AVRAM, P.

AVRAM, S.

Bukashet, "Mitsia d'Obiectivile si Creierimile Politice", Vol VI, No. 2, 1962	
<i>(Handwritten circled numbers 12, 27, and 28 are placed over the list)</i>	
1.	*Report on the Completion of Collectivization and the Reorganization of Agriculture Submitted to the Extra- ordinary Session of the <u>Trans National Assembly</u> at the Session of 27 April 1962, <u>Ed. ONUCITRO-DSS</u> ; pp 3-20.
2.	*The 014 Jordon Transmigration Network in Western Part of the Country Carried Out in Various Phasing Stages and the Functionality of the Migration in the State. <u>Report by Mr. A. TIRMAN</u> in <u>AJURA</u> and <u>MR. LILIAN</u> of the <u>Central Organization of Territory Central Organizational Territoriality</u> , pp 33-44.
3.	*The New M.P.B. 21/1958 Photocell Camera in Production. <u>Dr. MARCO</u> Candidate in Science (Candidate in Science) and <u>DR. V. ZHURKOV</u> ; pp 15-22.
4.	*The Necessity of Accepting the Plan for the Classification of Territory. <u>Ing. M. AVASTIĆ</u> and <u>Ing. I. CUREĆ</u> ; pp 55-59.
5.	*Program for the Appropriate Organization of the Territory in Support of the Socialist Agricultural Units, <u>Ing. A. ĐORĐEVIĆ</u> ; pp 60-63.
6.	*The Avram Institute, <u>Ing. S. AVRAM</u> , Center for the Orga- nization of Territory, Brochure; pp 1-83.

2000-0000-0000

— 2/ —

NEGREA, A.; BOTOSANEANU, I.; NEGREA, St. (Banat); TABACARU, I.; SERBAN, E.;
DANCAU, D.; AVRAM, S. (Cernisoara); DECU, A. (Oltenia); DECU, V.
(Oltenia); DUMITRESCU, M.; ORGHIDAN, T. (Cheile Virghisului);
TANASACHE, J.; GEORGESCU, M. (Dobrogea)

Contribution to the study of mollusks in Rumanian caves. Pt. 3.
Studii cerc biol anim 15 no.3:333-342 '63.

1. Comunicare prezentata de M.A. Ionescu, membru corespondent al
Academiei R.P.R.

AVRAM, Stefaniu

Observations on the variability of the species *Ischyropsalis dacica* Roewer - Rumanian ... (Ord. Opiliones, Subord. Palpatores Fam. Ischyropsalidae). Studii cerc. biol. s. zool. 16 no. 4:295-306 '64.

1. "Mihai Viteazul" Institute of Speleology.

AVP Stefania

A new opilionine, *Platybunus cirdeii* n. sp. Studii cerc biol s. zool
16 no. 6: 481-487 '64.

1. "Emil Racovita" Institute of Speleology.

"APPROVED FOR RELEASE: 06/06/2000

CIA-RDP86-00513R000102620004-6

HORZ, A.; SARO, E., chim.; ISTVAN, Fr.; BRADESCU, I.; ILIE, st.; URSESCU, I.
ing.; SAMI, Gh.; AVRAM, V., ing.

Drying some materials in fluidized bed. Rev chemic Min petr 1.5
no.7:409-411 J1 '64

APPROVED FOR RELEASE: 06/06/2000

CIA-RDP86-00513R000102620004-6"

AVRAMCHIK, M.N.

Growth of reindeer moss in the central part of the Anadyr River
Basin. Bot.shnr. 39 no.3:437-441 My-Ju '54. (MIRA 7:7)

1. Anadyrskaya optynaya olenevodcheskaya stantsiya.
(Anadyr Valley--Lichens) (Lichens--Anadyr Valley)

TSAGOLOV, N.A., prof., doktor ekon.nauk; BLYUMIN, I.G., prof., doktor ekon.nauk [deceased]; RUMYANTSEV, A.M., prof.; KORNIYENKO, A.A., dotsent, kand.ekon.nauk; SHNEIDERSON, A.I., prof., doktor ekon.nauk; LIP, Sh.B., prof., doktor ekon.nauk; SHVEDKOVA, G.M., kand.ekon. nauk; FISHEVSKIY, Yu.K.; DVORKIN, I.N., doktor ekon.nauk; SIDOROV, I.P.; KHAFIZOV, R.Kh., kand.ekon.nauk; NIKOLAEV, A.B., kand.ekon. nauk; AVRAMCHUK, F.P., kand.ekon.nauk; AL'TER, L.B., doktor ekon. nauk; BOYARSKIY, A.Ya., prof., doktor ekon.nauk; BREGEL', B.Ya., prof., doktor ekon.nauk; ARZUMANYAN, A.A.; VOLODIN, V.S., dotsent, kand.ekon.nauk; MIKSHA, L.S., kand.ekon.nauk; BUNKINA, M.K., dotsent, kand.ekon.nauk; YEVREYSKOV, A.V., kand.ekon.nauk; FADEYEVA, T.A., kand.ekon.nauk; KOLGANOV, M.V., prof., doktor ekon.nauk; KHRIMUSHIN, G.B., kand.ekon.nauk; MOSHENSKIY, M.G., kand.ekon.nauk; IVANOV, N.N., kand.ekon.nauk; GUTTSAYT, M.G., dotsent, kand.ekon. nauk; ABOLTIN, V.Ya., prof., doktor ekon.nauk; KOLLONTAY, V.M., kand.ekon.nauk; GLUKHAREV, L.I., kand.ekon.nauk; POKROVSKIY, A.I., kand.ekon.nauk; DADASHEV, G.A., dotsent, kand.ekon.nauk; ALESHPINA, I.V., kand.ekon.nauk; ZHANIN, V.A., dotsent, kand.ekon.nauk;

(Continued on next card)

TSAGOLOV, N.A.--(continued) Card 2.

KOZLOV, A.P.; TIMOFEEV, T.T., kand.istor.nauk; ALEKSEYEV, A.M.,
dotsent, kand.ekon.nauk; FILATOVA, Ye.M., dotsent, kand.ekon.nauk.
Prinimali uchastiyet: VOLKOV, F.M., kand.ekon.nauk; KHROMUSHIN,
G.B.; VOZNESENSKIY, L.A., nauchnyy sotrudnik. SPERANSKAYA, L., red.;
CHERKALEVA, C., tekhn.red.

[Criticism of present-day bourgeois, reformist, and revisionist
economic theories] Kritika sovremennoykh burzhuaznykh, reformistskikh
i revisionistskikh ekonomicheskikh teoriy. Pod red. N.A.Tsagalova.
Moskva, Izd-vo Sotsial'no-ekon.lit-ry, 1960. 588 p. (MIRA 13:5)

1. Moscow. Universitat. 2. Chlen-korrespondent AN SSSR (for Arsu-
manyam).

(Economics)

AVRAMCHUK, Fedor Petrovich, kand.ekonom. nauk; VLAS'YEVICH, Yuriy
Yevlampiyevich, kand. ekon. nauk; BEZDENEZHNYKH, P.T., red.;
CHAPAYEVA, N.I., tekhn. red.

[Militarization of the national economy and intensification
of the contradictions of imperialism] Militarizatsiia ekono-
miki i obostrenie protivorechii imperializma. Moskva,
Voenizdat, 196?. 141 p. (MIRA 15:10)
(Munitions) (World politics)

NEKRYACH, Ye. F.; SAMCHENKO, Z. A.; Prinimala uchastiye AVRAMCHUK, L. P.

Sorption of water vapor by hydrophilic high polymers. Part 8:
Isotherms of sorption and of heat of wetting of polycaprolactam.
Ukr. khim. zhur. 28 no. 5:514-621 '62. (MIRA 15:10)

(Nylon) (Sorption) (Heat of wetting)

NEKRYACH, Ye. F.; SAMCHENKO, Z. A.; Prinimala uchastiyu AVRAMOVU, L. P.

Sorption of water vapors by hydrophilic high polymers. Part 9:
Investigation of the structural changes of polycaprolactam based
on sorption and thermochemical data. Ukr. Khim. zhur. 28 no. 6:
703-706 '62. (MIRA 15:10)

1. Institut obshchey i neorganicheskoy khimi AN UkrSSR.

(Nylon) (Sorption) (Thermochemistry)

ACCESSION NR: AP4040772

S/0021/64/000/006/1774/0776

AUTHOR: Nekryach, Ye. F., Avramchuk, L. P.

TITLE: Kinetics of dissolution of synthetic polyamides [nylon, etc]

SOURCE: AN UkrSSR. Dopovidi, no. 6, 1964, 774-776

TOPIC TAGS: caprone, enant, yenant, anide, polyamide, polyamide dissolution, polyamide dissolution kinetics, methylene group, formic acid

ABSTRACT: The kinetics of the solution of polycaprolactam (caprone), polyhexamethylenedipamide (anide), and polyananthamide (enant) were investigated in aqueous solutions of formic acid of various concentrations at 20 °C. The study was motivated by the inadequacy of existing studies of the kinetics of solubility of polyamides. It was established that the solubility of polyamides depends to a great extent on their chemical structure, decreasing with an increase in the number of methylene groups in the elementary link of macromolecules of polymers. For 2-3 hours after the onset of dissolution, the percentage of dissolved polyamides η (%) is found to follow the empirical relation:

$$\lg \eta = \gamma \lg t + \lg B,$$

Where t is the time in solution, and γ and B are constants. Typical values for γ

Card 1/3

ACCESSION NR: AP4040772

and B for various concentrations of formic acid are given in Table 1 of Encl. 01.
Orig. art. has 5 graphs and one table.

ASSOCIATION: Instytut zagal'ny i neorganichny khimii AN UkrSSR (Institute of
General and Inorganic Chemistry, AN UkrSSR)

SUBMITTED: 25Sep63

ENCL: 01

SUB CODE: OC

NO REF SOV: 001

OTHER: 000

Card 2/3

ACCESSION NR: AP4040772

ENCLOSURE: 01

(a)	Monomer	Koeffizient des HCOOH, MOLE %	v	wB	w
(b)	Kempow	65,4	0,91	2,25	168
		63,1	0,91	2,30	170
(c)	Anilin	65,4	1,01	2,39	174
		63,1	1,01	2,50	178
(d)	Ehant	66,4	0,91	2,46	176

Table 1: Values of the coefficients
in the kinetic solubility equation

- (a) Polyanamide
- (b) Caprone
- (c) Anide
- (d) Yenant (enant?)
- (e) Concentration of formic acid (HCOOH), MOLES

Card 3/3

DUMANSKIY, A.V.; AVRMCHUK, L.P.; KURILENKO, O.D.; NEKRYACH, Ye.F.

Heat of reactions between a sulfonated styrene cationite and
water. Dokl. AN SSSR 159 no.5:1120-1122 D '64 (MIRA 181)

1. Institut obozrieni i neorganicheskoy khimii AN SSSR. 2. Chlen
korrespondent AN SSSR (for Dumanskiy).

ACC NR: AP7010716

SOURCE CODE: UR/0020/66.171/006/1373/1375

AUTHOR: Nekryach, Ye. F.; Gorkhovatskaya, N. V.; Avramchuk, L. P.; Kurilenko, O. D.; Dumanskiy, A. V. (Corresponding Member AN SSSR)

ORG: Institute of General and Inorganic Chemistry, Academy of Sciences Ukrainian SSR (Institut obshchey i neorganicheskoy khimii AN UkrSSR)

TITLE: Nature of exchange ions and the hydration energy of ionites

SOURCE: AN SSSR. Doklady, v. 171, no. 6, 1966, 1373-1375

TOPIC TAGS: ion exchange, heat of hydration, ionite

SUB CODE: 07

ABSTRACT: The authors state that while studying the heats of hydration of some hydrophilic polymers, they used ionites as a convenient model object for investigation. When wetting with water dry and moistened samples of K^+ , Na^+ , Ca^{2+} and Fe^{3+} forms of the sulfostyrene cationite KU-2 with a nominal divinylbenzene content of 4 and 20%, the heats increased in all cases in the order $K^+ < Na^+ < Ca^{2+} < Fe^{3+}$. This gave rise to the thought that there is a certain relationship between the energy of hydration and the charge of the counter ions. To check this supposition, the authors undertook to investigate the heats of wetting with water at 20° sulfo-

Card 1/2

UDC: 536.664 + 541.183.12

0930

3708

ACC NR: AP7010716

styrene cationite samples with the following exchange ions: single-charged Cs⁺, Rb⁺, K⁺, Na⁺, Li⁺; doubly-charged Ba²⁺, Ca²⁺, Mg²⁺; and triply-charged Fe³⁺, Al³⁺. At the same time, water-vapor sorption isotherms were taken for the same samples at 20° on a vacuum sorption apparatus. The authors state that the results justify the assertion that a direct relationship exists between the size of the charge of exchange ions and the hydration energy of ionites as determined from the heats of wetting them with water. Orig. art. has: 1 figure. [JPRS: 40,351]

Card 2/2

L 2106-66 EWT(1)/EWP(m)/FS(v)-3/EWA(d) 0W

ACCESSION NR: AP5024779

UR/0021/65/000/009/1139/1145

AUTHOR: Klikh, Yu. O. (Klikh, Yu. A.); Avramchuk, N. A.

TITLE: Certain cases of the perturbed motion of a mass point in a central gravitational field

SOURCE: AN UkrRSR. Dopovidi, no. 9, 1965, 1139-1145

TOPIC TAGS: spacecraft motion, spacecraft trajectory, asymptotic method, nonlinear mechanics

ABSTRACT: The Krylov-Bogolyubov method is applied to solving the problem of motion of a flight vehicle with a small constant thrust in a central gravitational field. The flight vehicle is considered as a point whose mass m varies linearly. The equations of motion are transformed to a form with a small parameter ϵ ($\epsilon = \dot{m}/m_0$, where m_0 is the initial mass of m and \dot{m} is the rate of expenditure of the mass) multiplying the nonlinear terms. This form of the equations makes it possible to apply asymptotic methods of nonlinear mechanics to the solution of the problem. By applying the averaging method, equations of the first approximation are derived which are valid for any direction of the thrust force. The equations of the first approximation are integrated and the trajectories of motion are determined for the

Card 1/2

L 2106-66

ACCESSION NR: AP5024779

following two particular cases: a) radial thrust, and b) constant direction thrust. Orig. art. has: 1 figure and 23 formulas. [LK]

ASSOCIATION: Odes'kyy politekhnichnyy instytut (Odessa Polytechnic Institute)

44

SUBMITTED: 26Sep64

ENCL: 00

SUB CODE: SV, ME

NO REF SOV: 002

OTHER: 004

ATD PRESS: 4113

Cord 2/2

LUGOVSKIY, Sergey Ivanovich; DYMCHUK, Gennadiy Konstantinovich;
DROBOT, Boris Yakovlevich; AVRANCHUK, Rostislav Nikiforovich.
Prinimali uchastiye: MAR'YENKOV, V.V.; BAKIROV, U.Kh.;
NIKITIN, V.S., kand. tekhn. nauk, retsenzent; STEBAKOV, B.A.,
gorn. inzh., otv. red.

[Ventilation of mines and strip mines] Ventiliatsiya shakht i
kar'erov. [By] S.I.Lugovskii i dr. Moskva, Izd-vo "Nedra,"
1964. 306 p. (MIRA 17:5)

AVRAMCHUK, R.N., inzh.

Ventilating mines during multi-level working of unconnected deposits. Sbor. nauch. trud. MGRI no.10:80-8 '61
(MIRA 17:8)

Efficient method of ventilating developed horizons during multi-level mining. Ibid. 88-92

L 24804-65 EWG(v)/EMT(1)/EEG(t)
ACCESSION NR: AT4049980

Po-4/Pe-5/Pac-2 GW/HLK
S/0000/04/000/000/0003/0015

AUTHOR: Avramchuk, V. V.

TITLE: Polychromatic polarization of some lunar areas ✓

SOURCE: AN UkrSSR. Glavnaya astronomicheskaya observatoriya. Fizika Luny i planet
(Physics of the moon and planets). Kiev, Naukova dumka, 1964, 3-16

TOPIC TAGS: moon, lunar polarization, Umov effect, polychromatic polarization

ABSTRACT: The author gives a brief outline of the meaning of the Umov effect as interpreted by G. V. Rozenberg (UFN, 19, No. 1, 57, 1959). It is postulated that in the case of the moon the dependence of polarization on wavelength is due to some other factor than the Umov effect. The author gives the results of polarization measurements of the moon in 18 cross sections of the spectrum over a range of 355 - 600 nm. The observations were carried out in September 1962. An analysis of these data shows that at a phase of +12 - 13° the polarization of the investigated details increases toward the red end of the spectrum, which contradicts the results obtained by E. K. Cohan. For other phases the degree of polarization is an almost monotonically varying function of wavelength which increases intensely in ultraviolet light ($\lambda_{eff} = 358$ nm). The author has also established a dependence between the

Cord 1/2

L 24804-65

ACCESSION NR: AT4049980

maximum degree of polarization P_{\max} and albedo A in narrow parts of the spectrum ($\lambda_{eff} = 430$ and 560 nm). At the phase -20° , polarization was not detected in any part of the spectrum investigated. Comparison with the results obtained by E. K. Cohan and K. N. Derkach (Tr. Khar'k. astron. obs., 14, 79, 1962) shows that the results of the latter are 40% below the true values. "In conclusion, the author wishes to thank A. V. Murkov and V. P. Kompleya who were responsible for the formulation of this study." Orig. art. has: 6 formulas, 6 figures and 2 tables.

ASSOCIATION: None

SUBMITTED: 07May64

ENCL: 00

SUB CODE: AA

NO REF SOV: 012

OTHER: 004

Cord 2/3

L 3219-65 EWT(1)/FCC GS/GW
ACCESSION NR: AT5024608

UR/0000/65/000/000/0112/0120

AUTHOR: Avramchuk, V. V.

TITLE: Multicolor polarimetry of the light of the twilight and day sky at the zenith

SOURCE: AN UkrSSR. Voprosy astrofiziki; issledovaniye atmosfer Venery i Marsa (Problems in astrophysics; investigation of the atmospheres of Venus and Mars). Kiev, Izd-vo Naukova dumka, 1965, 112-120

TOPIC TAGS: twilight, day sky, polarimetric investigation, light polarization, planetary atmosphere, atmospheric optics

ABSTRACT: Measurements of the degree of light polarization of the twilight and day sky in 12 spectral regions of the 355—600 nm band were carried out with the 70-cm Cassegrainian reflector and electropolarimeter with rotating polaroid of the Main Astronomical Observatory of the Ukrainian Academy of Sciences. The measurements show that the observed values of the degree of polarization in the zenith are considerably less for all wavelengths than the theoretically derived values of polarization of first-order scattering computed for a Rayleigh atmosphere. Data show further that the value of maximum polarization never exceeds

L 3219-66

ACCESSION NR: AT5024608

70-75% for the red region of the spectrum and 50-60% for the ultra-violet. The results of observations indicate that in the spectral interval 420-600 nm the degree of polarization has only a negligible dependence on wavelength. A sharp drop in the degree of polarization is evident at 420 nm. Orig. art. has 4 figures and 6 tables. [DM]

ASSOCIATION: Glavnaya astronomicheskaya observatoriya AN UkrSSR (Main Astronomical Observatory, AN UkrSSR)

SUBMITTED: 05Jun65

ENCL: 00

SUB CODE: ES

NO REF Sov: 004

OTHER: 000

ATD PRESS: 4104

PC

Copy 2/2

PAVLOV, I.F., kand.biolg.nauk; LEBEDEVA, K.K., nauchnyy sotrudnik;
AVRAMENKO, A.I., starshiy tekhnik

Methods for protecting grain crops. Zashch. rast. ot vred.
i bol. 7 no.7:22-24 Jl '62. (MIRA 15:11)

(Central Black Earth Region—Grain—Diseases and pests)
(Central Black Earth Region—Plants, Protection of)

ACC NR: AP6015710

(A)

SOURCE CODE: UR/0413/66/000/009/0125/0125

INVENTOR: Naydis, N. M.; Avramenko, A. K.; Yakuts, B. L.; Ryzhov, L. S.; Korchin, Yu. M.; Kalyuzhnyy, O. K.; Kuchinskiy, V. A.

ORG: None

TITLE: Fuel delivery controller for internal combustion engines. Class 46, No. 181445

SOURCE: Izobreteniya, promyshlennyye obraztsy, tovarnyye znaki, no. 9, 1966, 125

TOPIC TAGS: engine fuel system, air temperature, fuel control

ABSTRACT: This Author's Certificate introduces: 1. A fuel delivery controller for internal combustion engines. The unit consists of a device for transmitting signals to a servomechanism, a stack of aneroid capsules and two correctors with pickups. These pickups are made in the form of bimetallic plates equipped with manual adjustment screws. Each of these bimetals varies fuel delivery as a function of air temperature. The second corrector is connected to the fuel delivery channel supplying fuel to the engine to allow for the variation in the specific weight of the fuel with temperature. 2. A modification of this controller in which transition from one type of fuel to another is simplified by a scale on the device for correcting temperature (specific weight). The indicating needle of the corrector scale can be set by a manual adjustment screw.

SUB CODE: 21 / SUBM DATE: 28Jun63

UTC: 621.43,031-441.2

Card 1/1

AKHUNDOV, E.B.; AVER'YENKO, A.V.; BAMPI, Yu.S.

Optimum power of a condensing electric power plant
operated on peat. Trudy Inst.energ. AN BSSR no.10:12-21
'59. (Peat) (Electric power plants)

(MIRA 13:6)

TROITSKIY, N.A.; AVRAMENTO, B.I.

Correlation between the dose and absorbed dose in the irradiation of
seeds. Radiobiologija 4 no.1:180-181 '64. (MIRA 17:4)

1. Institut biologii AN BSSR, Minsk.

L 55133-6
ACCESSION NR: AP501000

UR/0250/65/009/003/0202/0204 .2/

AUTHOR: Avramenko, B. I.; Ipat'yev, A. N.; Mushinskaya, L. G.; Savchenko, A. V.; Zhebrak, A. R.

TITLE: Male sterility in plants caused by penetrating radiation

SOURCE: AN BSSR. Doklad, v. 9, no. 3, 1965, 202-204

TOPIC TAGS: radiobiology, gamma ray, cobalt 60, radiation effect, seed, plant genetics

ABSTRACT: It is a laborious and costly process to obtain hybrid seeds by the usual method of flower castration. The authors studied the possibility of inducing male sterility in plants by irradiating air-dried cucumber, rye, wheat, tomato, radish and other seeds with gamma rays from Co⁶⁰ in the atomic reactor of the AN BSSR. Critical doses for each species of plants were used. A relationship was noted between sterility and the radiation dose in the case of mustard, cabbage, cucumber, and rape. In corn and beans, some varieties were less sensitive than others to the same radiation dose. Pollen was found to be sterile in non-irradiated plants, indicating that male sterility may be due to some other factors. In

Card 1/2

L 55133-65

ACCESSION NR: AP5011090

general, however, the results of the experiments showed that irradiation of seeds increases pollen sterility so that joint planting of an irradiated maternal variety with a non-irradiated paternal variety increases the hybridity of the seeds. Orig. art. has: 4 tables.

ASSOCIATION: Otdel genetiki i tsitologii AN BSSR (Genetics and Cytology Section AN BSSR)

SUBMITTED: 29Jan64

ENCL: 00

SUB CODE: LS

NO REF SOV: 017

OTHER: 003

Card 2/2

AVRAMENKO, V.D.

For a fundamental change of the system of standardizing the quality
of coal. Ugol' 30 no. 216-9 P '55. (MLRA 8:4)

1. VNIUgleobogashcheniye.
(Coal)

AVRAMENKO, B.I.; IPAT'YEV, A.N.; MUSHINSKAYA, L.G.; SAVCHENKO, A.P.

Morphological and biological changes in plants subjected to
gamma irradiation. Dokl. AN BSSR 9 no. 5:340-343 My '65
(MIRA 1981)

1. Institut genetiki i tsitologii AN BSSR. Submitted February
28, 1964.

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ACC NR: AP6014957

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35

AUTHOR: Avramenko, B. I.; Ipt'yev, A. N.; Mushinskaya, L. G.; Savchenko, A. P.

B

ORG: Institute of Genetics and Cytology, AN BSSR (Institut genetiki i tsitologii
AN BSSR)

TITLE: Morphological and biological changes in plants induced by gamma rays

19

SOURCE: AN BSSR. Doklady, v. 9, no. 5, 1965, 340-343

TOPIC TAGS: gamma ray, radiation plant effect, plant chemistry

ABSTRACT: Critical and sublethal doses of gamma rays stunted the growth of tomatoes, cucumbers, cabbage, mustard, radishes, beans, beets, and onions. Seeds exposed to such doses germinated 1-14 days later than did the control. Subsequent development was also slower. These doses likewise altered the plants' morphology, particularly the leaves. However, all the changes gradually disappeared by the time the plants flowered, indicating that plants recover at a certain stage of development, even after receiving very high doses of radiation. Irradiation also affected the biochemical composition of the plants. For example, it reduced the fat content of mustard and cabbage seeds below that of the control.

Low doses of gamma rays, on the other hand, had a stimulating effect. They hastened the ripening of the fruits and increased the plants' productivity. This paper was presented by Academician AN BSSR A. R. Zhebrak. Orig. art. has: 3 tables. [JPRS]

SUB CODE: 06 / SUBM DATE: 28Feb64 / ORIG REF: 007 / OTH REF: 003

Card 1/1 B1K

2

8(6)

PHASE I BOOK EXPLOITATION

SOV/2382

Avramenko, F.D., V.I. Veyts, B.A. Gurevich, V.I. Denisov, A.G. Zakharin,
N.A. Karaulov, I.S. Kolosov, N.N. Krachkovskiy, S.N. Kritskiy, M.M.
Lebedev, T.K. Leont'yeva, M.F. Menkel', A.S. Nekrasov, G.I. Rossiyevskiy,
and B.I. Shvorin

Osnovnyye voprosy planirovaniya yedinoy energeticheskoy sistemy SSSR (Basic
Problems in Planning a Unified Power System for the USSR.) Moscow,
Izd-vo AN SSSR, 1959. 174 p. Errata slip inserted. 2,500 copies printed.

Sponsoring Agency: Akademiya nauk SSSR. Energeticheskiy institut.

Eds.: G.M. Krzhizhanovskiy, Academician and V.I. Veyts, Corresponding
Member, USSR Academy of Sciences; Tech. Ed.: S.G. Markovich.

PURPOSE: This book is intended for government planning circles, scientific
research organizations and others interested in the electrification of the
USSR.

COVERAGE: The book examines the principal problems of a unified power system

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Basic Problems (Cont.)

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of the system, the increasing use of semiconductors, the use of various types of fuels, etc. These problems were presented in two earlier publications of the Academy of Sciences: Nauchnyye osnovy sozdaniya i razvitiya yedinoj energeticheskoy sistemy SSSR (Scientific Bases in the Creation and Development of a Unified Power System in the USSR; Conclusions of a Coordinating Conference, Moscow, 1957); and Razrabotka nauchnykh osnov razvitiya energeticheskikh sistem i ikh ob'yedineniya yedinoj energeticheskoy sistemy

(Working Out of Scientific Bases in the Development of Power Systems and Their Integration Into a Unified Power System. Series: Voprosy sovetskoy nauki, Moscow, 1958). The following persons participated in writing the book: F.D. Avramenko (Chapters 2 and 4); V.I. Veyts (Chapters 2, section 4 of Chapter 3, Chapter 4, section 1 of Chapter 6, Chapters 8 and 9); B.A. Gurevich (Chapter 1, section 1 of Chapter 7); V.I. Denisov (Chapters 4 and 8); A.G. Zakharin (section 2 of Chapter 7); N.A. Karaulov, S.N. Kritskiy and M.F. Menkel' (Chapter 5); N.N. Krachkovskiy (section 4 and 5 of Chapter 6); I.S. Koslov (section 8 of Chapter 1); M.M. Lebedev (Chapter 6, section 1 of Chapter 7, Chapters 9,10,11); T.K. Leont'yeva (section 1 of Chapter 3); A.S. Nekrasov (sections 2 and 3 of Chapter 9); G.I. Rossiyevskiy (Chapter 3); B.I. Shvorin (Chapter 2). Those who participated in preparing the material were: M.M. Albegov, K.N. Bestuzheva, V.A. Bondareva, M.S. Vdovchenko, A.L. Valikanov, Ye.A. Volkova, V.A. Gadiyeva, I.U. Kon'ya, D.N. Korobova, Yu.S. Kretinina, M.A.

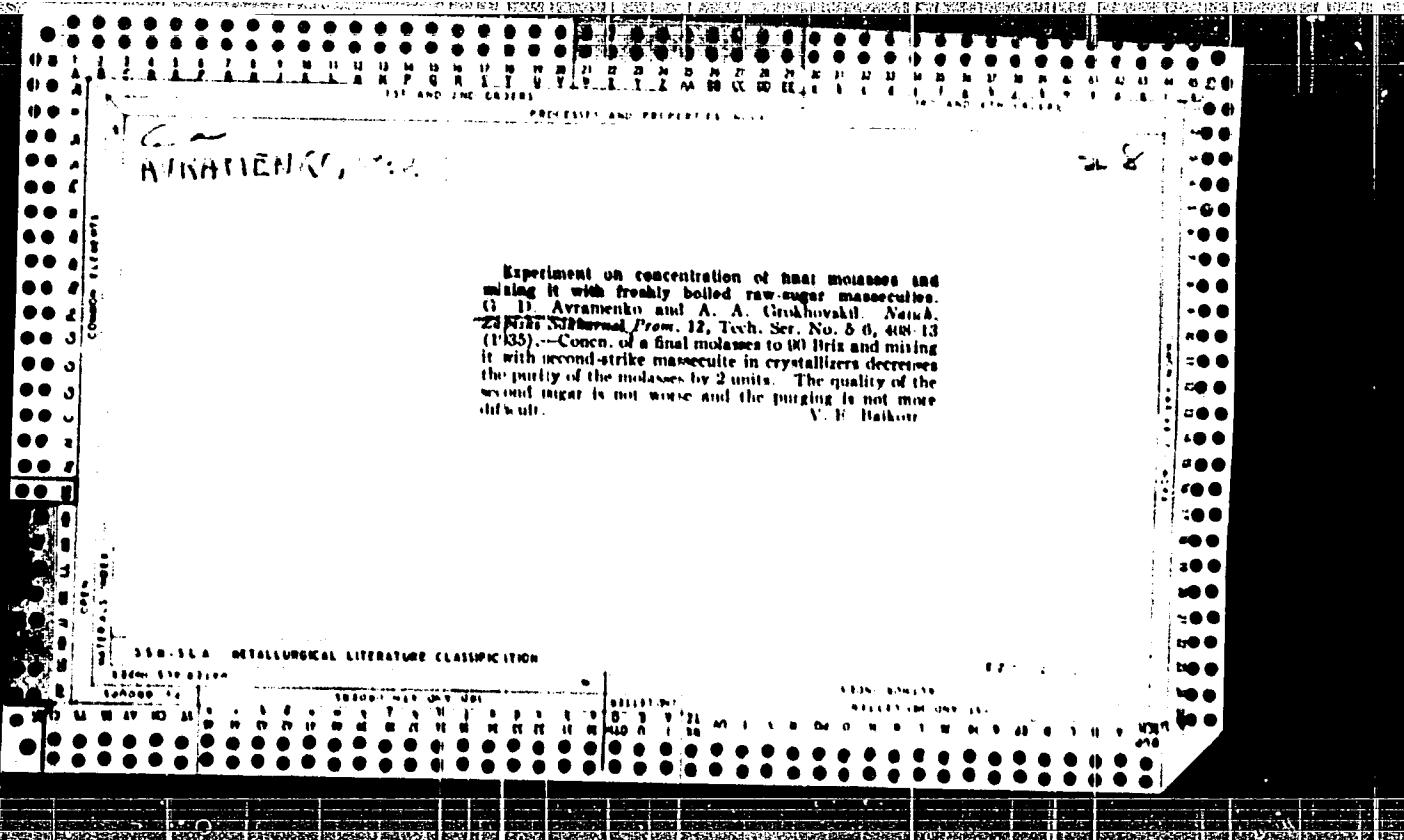
Card 3/11

AVRAMENKO, Fedor Dmitriyevich; BRUSANOVSKIY, Semen Genrikhovich;
MAKAROVA, E.A., red.; DROZDOV, G.M., tekhn. red.

[Magnitogorsk and Kuznetsk; competition between the steelworkers
of the Urals and Siberia] Magnitka-Kuznetsk; sorevnovanie metal-
lurgov Urala i Sibiri. Moskva, Izd-vo VTsSPS Profizdat, 1962.
97 p.

(MIRA 15:5)

(Novokuznetsk--Steel industry)
(Magnitogorsk--Steel industry)
(Socialist competition)



CA R/ANTHELIC, 6/1

-38

Improvement of vacuum pan feed line. G. D. Avra-
menko. Tzidhermaya Prom. 22, No. 7, 42-3(1948).—
The feed pipe must be installed in the center of the intake
and end about 30 cm. above the discharge valve.
V. K. Balkow

ASSISTANT METALLURGICAL LITERATURE CLASSIFICATION

BOOK SHELVING

BOOK SHELVING

BOOK SHELVING	BOOK SHELVING
BOOK SHELVING	BOOK SHELVING

AVRAMENKO, G. I.

37434. Opyt razdoya korov v kolkhozakh molokovskogo zoovetuchastka, Leninsky
rayon mosk. Obl/ Sets. zhivotnovodstvo, 1949, No. 8, s. 40-45.

SO: Letopis' Zhurnal'nykh Statey, Vol. 7, 1949

AVRAMENKO, G. I. Cand Agr Sci -- (diss) "Effect of various types of combination of succulent fodders upon the growth and development of heifers and their subsequent milk productivity." Mos, 1959. 17 pp (All-Union Order of Lenin Acad Agr Sci im V. I. Lenin. All-Union Sci Res Inst of Animal Husbandry), 150 copies (KL, 45-59, 148)

AVRAMENKO, I., redaktor; KIRILLINA, L., tekhnicheskiy redaktor.

[Youth outstrips time; the participation of Communist Youth and young people in the struggle to raise labor productivity] Molodezh' obgoniaet vremia; ob uchastii komsomol'tsev i molodezhi v bor'be za povyshenie proizvodstvennoi truda. Moskva, "Molodaiia gvardiia," 1955. 79 p.

(Labor productivity)

(MLRA 8:5)

KUDRYAVTSEVA-MOLODCHIKOVA, Larisa Pavlovna; AVRAMENKO, I., red.;
KHAKHIN, M., tekhn, red.

[Grain ears; stories from the life of cultivated cereals]
Khlebnye kolos'ia; rasskazy iz zhizni kul'turnykh zlakov.
Moskva, Izd-vo TAK VREM "Molodaia gvardiia," 1959. 174 p.

(Grain)

(MIRA 12:10)

KUDRAVTSEVA-MOLOCHIKOVA, Larisa Pavlovna; AVRAMENKO, I., red.;
VOLKOVA, L., red.

[History of an apple trees; true stories on the wonderful
life of a fruit tree] Istoriia odnoi jablon'ki; pravdivye
rasskazy ob udivitel'noi zhizni plodovogo dereva. Moskva,
Molodaia gvardiia, 1964. 126 p. (MIRA 17:4)

RUDNEV, D.F.; AVRAMENKO, I.D.; KRAZHEVSKAYA O.N.

Using DDT oil solutions for combating pests in the Kiev greenbelt.
Nauch. trudy Inst. ent. i fit. 6:80-88 '55. (MIRA 9:?)
(Kiev--DDT (Insecticide)) (Kiev--Trees--Diseases and pests)

AVRAMENKO, I. D., Cand Biol Sci (diss) -- "Bark-chewing rodents of the Kazan'-Veshenskaya massif and measures for combatting them". Khar'kov, 1960. 17 pp (Min Agric Ukr SSR, Khar'kov Order of Labor Red Banner Agric Inst im V. V Dokuchayev), 200 copies (KL, No 14, 1960, 129)

MARDZHANYAN, G.M.; ASATRYAN, E.V.; MARKOSYAN, A.A.; UST'YAN, A.K.;
AVRAMENKO, I.D., kand. biolog. nauk (Gomel'); MISKO, L.A.;
AGAFONOVA, Z.Ya., kand. biolog. nauk; ABBASOV, Ya.M., mladshiy
nauchnyy sotrudnik; SADYKHOV, D.M., aspirant

Brief information. Zashch. rast. ot vred. i bol. 8 no.10:
55-57 O '63. (MIRA 17:6)

1. Armyanskiy institut zemledeliya (for Mardzhanyan, Asatryan,
Markosyan, Ust'yan). 2. Poltavskiy sel'skokhozyaystvennyy
institut (for Misko). 3. Kurskaya sel'skokhozyaystvennaya
opytnaya stantsiya (for Agafonova). 4. Azerbaydzhanskiy
nauchno-issledovatel'skiy institut khlopkovodstva, Kirovabad
(for Abbasov). 5. Vsesoyuznyy institut zashchity rasteniy (for
Sadykhov).

USSR/Diseases of Farm Animals. Noninfectious Diseases R-2

Abs Jour : Ref Zhur-Biol., No 2, 1958, 2766

Author : Avramenko I. F.

Inst : Not given

Title : Cause of Diarrhea in Newborn Lambs

Orig Pub : Oytevoda, 1957, No 4, 42-44

Abstract : Clinical observations and experimental tests have shown that poor care and low food value of the fodder given to pregnant ewes and newborn lambs are the reasons for the mass incidence of diarrhea in lambs and their high mortality (up to 50%) in kolkhozes in a number of rayons in Buryat-Mongolian ASSR. By improving the care and fodder given to pregnant ewes, and the administration of sintomycin or biomycin to newborn lambs it was possible to prevent the lambs from contracting

Card 1/1

VASILENKO, S.I.; MEZHAKOV, V.A.; AVRAMENKO, I.I.; ASTAKHOV, A.V., otvetstvennyy
red.; SILINA, L.A., red.; ALDANOVA, Ye.I., tekhn. red.

["Shakhter-2" coal cutter-loader] Ugol'nyi kombain "Shakhter-2"
Moskva, Ugletekhnizdat, 1958. 147 p. (MIRA 11:7)
(Coal mining machinery)

KSENZUK, F.A., inzh.; AVRAMENKO, I.N., inzh.; MIRENSKIY, Yu.M.; TROSHCHENKOV,
N.A.

Relation between the degree of deformation and the speed and tension
during the straightening of sheet steel for automobiles. Stal' 25
no.7:632-634 JI '65. (MIRA 18:7)

1. Zavod "Zaporozhstal".

ACC NR: AT6012089

(N)

SOURCE CODE: UR/3177/65/021/000/0038/0052

AUTHOR: Chekmarev, A. P. (Academician AN UkrSSR); Suf'yan, M. M. (Professor);
Meleshko, V. I. (Candidate of technical sciences); Prokof'yev, V. I. (Candidate of technical
sciences); Avramenko, I. N. (Engineer); Dodoka, V. G. (Engineer); Ksenzuk, F. A. (Engineer);
Kudin, D. P. (Engineer); Lola, V. N. (Engineer); Movshovich, V. S. (Engineer); Pavlishchev,
V. B. (Engineer); Soroko, L. N. (Engineer); Sukhobrus, Ye. P. (Engineer); Kholodnyy, V. P.
(Engineer); Yudin, M. I. (Engineer)

ORG: none *

TITLE: Improvements in the techniques of production of Kh18Ni0T cold-rolled wide-strip
steel at the Zaporozhstal' Plant

SOURCE: Dnepropetrovsk. Institut chernoy metallurgii. Trudy, v. 21, 1965. Prokatnoye
proizvodstvo (Welding production), 38-52

TOPIC TAGS: stainless steel, bright stock lubricant, metal rolling, sheet metal, industrial
plant / Kh18Ni0T stainless steel, P-28 bright stock lubricant

ABSTRACT: On increasing to 11.8 tons from the previous 10.3 tons the weight of the ingots

Cord 1/2

URNS, I.G.; AVRAENKO, K.I.

Cysts and tumors of the mediastinum. Trudy Vor. med. inst. 52.
221-225 '63. (SIRU 18.3)

AVRAMENKO, I. I.; KOLESNIKOVA, R. V.; KUZNETSOVA, N. I.

Reaction rate constants and the mechanism of reactions of oxygen atoms
with methane and ethane. Izv. AN SSSR. Otd. khim. nauk no. 4:620-627
Ap '63. (MI:A 15:3)

1. Institut khimicheskoy fiziki AN SSSR.
(Paraffins) (Oxygen) (Chemical reaction, Rate of)

KITLINSKAYA, Vera Kazimirovna; AVRAMENKO, I.K., red.; BHUSILOVSKAYA, S.I.,
tekhn. red.

[China today and tomorrow] Kitai segodnia i zavtra. Leningrad,
Sovetskii pisatel', 1958. 393 p.
(MIRA 11:8)
(China--Description and travel)